

Eylor
08/644289

08/644289

FILE 'REGISTRY' ENTERED AT 15:48:13 ON 24 JUN 1999

L12 1 S SLRPFKALVREKGHRPSHSC/SQSP

=> d .bevreg1

L12 ANSWER 1 OF 1 REGISTRY COPYRIGHT 1999 ACS

RN 173787-20-7 REGISTRY

CN L-Cysteine, L-seryl-L-leucyl-L-arginyl-L-prolyl-L-phenylalanyl-L-lysyl-L-alanyl-L-leucyl-L-valyl-L-arginyl-L-.alpha.-glutamyl-L-lysylglycyl-L-histidyl-L-arginyl-L-prolyl-L-seryl-L-histidyl-L-seryl-
(9CI) (CA INDEX NAME)

SQL 20

SEQ 1 SLRPFKALVR EKGHRPSHSC

=====

HITS AT: 1-20

REFERENCE 1: 128:216370

REFERENCE 2: 128:21873

REFERENCE 3: 124:173450

(FILE 'CAPLUS' ENTERED AT 15:49:12 ON 24 JUN 1999)

L13 3 S L12

=> d 1-3 .bevstr

L13 ANSWER 1 OF 3 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1998:178120 CAPLUS

DOCUMENT NUMBER: 128:216370

TITLE: Recombinant p53as protein and antibody for diagnosis and therapy of malignancy

INVENTOR(S): Kulesz-Martin, Molly F.

PATENT ASSIGNEE(S): Health Research, Inc., USA

SOURCE: U.S., 30 pp. Cont.-in-part of U.S. Ser. No. 259,612.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 6

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5726024	A	19980310	US 96-644291	19960510
US 5688918	A	19971118	US 94-259612	19940614
EP 806478	A2	19971112	EP 97-107696	19970512
EP 806478	A3	19990609		

Searcher : Shears 308-4994

R: BE, CH, DE, DK, FR, GB, LI, NL, SE

PRIORITY APPLN. INFO.: US 93-100496 19930802
 US 94-195952 19940211
 US 94-259612 19940614
 US 96-644289 19960510
 US 96-644291 19960510
 US 96-644456 19960510

AB The invention comprises plasmids and viral vectors contg. an animal p53as (alternatively spliced p53) cDNA sequence. A portion of the p53as sequence may be identified to a position of wild type p53 gene from the same animal. In preferred embodiments, the p53as is mouse or human p53as. A preferred viral vector is baculovirus vector. The invention further includes antibodies both polyclonal and monoclonal, to p53as and to at least a portion of human p53 intron 10 sequence encoding SLRPFKALVREKGRPSSHSC, which is related to p53as sequences and plasmids and viral vectors contg. such sequences. All of the above find utility in studying p53 and p53as and their relative expressions which is believed important for detection and control of malignant cells and their susceptibility to treatment agents. The antibodies can detect the presence of p53as and related sequences and when injected into cells could cause cell cycle arrest and the plasmids and viral vectors, with appropriate promoters, can cause expression of the p53as and p53 intron 10 sequences which can affect cell growth and perhaps arrest certain malignancies.

IT 173787-20-7

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (recombinant p53as protein and antibody for diagnosis and therapy of malignancy)

L13 ANSWER 2 OF 3 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1997:761610 CAPLUS
 DOCUMENT NUMBER: 128:21873
 TITLE: P53as protein and antibody therefor
 INVENTOR(S): Kulesz-Martin, Molly F.
 PATENT ASSIGNEE(S): Health Research, Inc., USA
 SOURCE: U.S., 25 pp. Cont.-in-part of U.S. Ser. No. 195,952.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 6
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5688918	A	19971118	US 94-259612	19940614
EP 652232	A1	19950510	EP 94-610042	19940801

Searcher : Shears 308-4994

08/644289

R: BE, CH, DE, DK, FR, GB, LI, NL, SE

JP 08099998	A2	19960416	JP 94-181558	19940802
CA 2150994	AA	19951215	CA 95-2150994	19950605
JP 08081500	A2	19960326	JP 95-169323	19950613
EP 709397	A1	19960501	EP 95-610034	19950614

R: BE, CH, DE, DK, FR, GB, LI, NL, SE

US 5726024	A	19980310	US 96-644291	19960510
------------	---	----------	--------------	----------

PRIORITY APPLN. INFO.:

US 93-100496	19930802
--------------	----------

US 94-195952	19940211
--------------	----------

US 94-259612	19940614
--------------	----------

AB The invention comprises plasmids and viral vectors contg. an animal p53as cDNA sequence. A portion of the p53as sequence may be identified to a position of wild type p53 gene from the same animal. In preferred embodiments, the p53as is mouse or human p53as. A preferred viral vector is baculovirus vector. The invention further includes antibodies both polyclonal and monoclonal, to p53as and to at least a portion of human p53 intron 10 sequence encoding SLRPFKALVREKGRPSHSC which is related to p53as sequences and plasmids and viral vectors contg. such sequences. All of the above find utility in studying p53 and p53as and their relative expressions which is believed important for detection and control of malignant cells and their susceptibility to treatment agents. The antibodies can detect the presence of p53as and related sequences and when injected into cells could cause cell cycle arrest and the plasmids and viral vectors, with appropriate promoters, can cause expression of the p53as and p53 intron 10 sequences which can affect cell growth and perhaps arrest certain malignancies.

IT 173787-20-7

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(recombinant p53as protein and antibody for diagnosis and therapy of malignancy)

L13 ANSWER 3 OF 3 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1996:126792 CAPLUS

DOCUMENT NUMBER: 124:173450

TITLE: Monoclonal and polyclonal antibody to alternatively spliced p53 protein (p53as) for diagnosis and prognosis of cancer

INVENTOR(S): Kulesz-Martin, Molly F.

PATENT ASSIGNEE(S): Health Research, Inc., USA

SOURCE: Can. Pat. Appl., 40 pp.

CODEN: CPXXEB

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 6

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	Searcher	:	Shears	308-4994

-----	-----	-----	-----	-----
CA 2150994	AA	19951215	CA 95-2150994	19950605
US 5688918	A	19971118	US 94-259612	19940614
PRIORITY APPLN. INFO.:			US 94-259612	19940614
			US 93-100496	19930802
			US 94-195952	19940211

AB The invention comprises plasmids and viral vectors contg. an animal p53as cDNA sequence. A portion of the p53 sequence may be identified to a position of wild type p53 gene from the same animal. In preferred embodiments, the p53as is mouse or human p53as. A preferred viral vector is baculovirus vector. The invention further includes antibodies both polyclonal and monoclonal, to p53as and to at least a portion of human p53 intron 10 sequence encoding SLRPFKALVREKGHRPSHSC which is related to p53as sequences and plasmids and viral vectors contg. such sequences. All of the above find utility in studying p53 and p53as and their relative expressions which is believed important for detection and control of malignant cells and their susceptibility to treatment agents. The antibodies can detect the presence of p53as and related sequences and when injected into cells could cause cell cycle arrest and the plasmids and viral vectors, with appropriate promoters, can cause expression of the p53as and p53 intron 10 sequences which can affect cell growth and perhaps arrest certain malignancies.

IT 173787-20-7

RL: BSU (Biological study, unclassified); BIOL (Biological study)
 (polyclonal or monoclonal antibody to human or mouse p53as
 proteins for diagnosis or prognosis of cancer)

=> fil hom

FILE 'HOME' ENTERED AT 15:49:47 ON 24 JUN 1999

 MWSEKELI
 (TM)

Release 3.1A John F. Collins, Biocomputing Research Unit.
 Copyright (c) 1993-1998 University of Edinburgh, U.K.
 Distribution rights by Oxford Molecular Ltd

Msearch_p protein - protein database search, using Smith-Waterman algorithm

Run on: Wed Jun 23 14:53:55 1999; Maspar time 41.91 Seconds

Tabular output not generated. 10.148 Million cell updates/sec

Title: >US-08-644-289-1

Description: (1-20) from US08644289.pep

Sequence: 1 SLRPFKALVREKGRHPSHC 20

Scoring table: PAM 150

GAP 15

Searched: 170751 seqs, 2126608 residues

Post-processing: Minimum Match 0%

Listing first 1000 summaries
 Maximum DB seq length 30

Database:

a:geneseqs
 1:part1.2:part2.3:part3.4:part4.5:part5.6:part6.7:part7
 8:part8.9:part9.10:part10.11:part11.12:part12.13:part13
 14:part14.15:part15.16:part16.17:part17.18:part18
 19:part19.20:part20.21:part21.22:part22.23:part23
 24:part24.25:part25.26:part26.27:part27.28:part28
 29:part29.30:part30.31:part31.32:part32.33:part33
 34:part34.35:part35.36:part36.37:part37.38:part38
 39:part39

Statistics: Mean 20.743; Variance 62.468; scale 0.332

Pred. No. is the number of results predicted by chance to have a
 score greater than or equal to the score of the result being printed,
 and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	Pred. No.
1	151	100.0	20	18	R92698 Human wild-type p53 g	5.67e-10
2	136	90.1	21	29	W33235 Human p53 fragment.	4.52e-08
3	85	56.3	20	29	W33239 Peptide of the human	7.01e-02
4	50	33.1	18	8	R44726 HEV ORF1 peptide (123	4.09e+02
5	48	31.8	17	29	W33236 Mouse p53as peptide.	6.37e+02
6	47	31.1	24	7	R37861 HIV protein fragment	7.92e+02
7	46	30.5	16	9	R45673 RNP heparin binding	9.83e+02
8	46	30.5	20	26	W26230 pMA2-TF-1g6 junction	9.83e+02
9	45	29.8	14	6	R45671 N terminus of human C	9.83e+02
10	45	29.8	14	6	R32978 Mastoparan analogue (1.22e+03
11	45	29.8	19	24	W28500 Last 19 C-terminal am	1.22e+03
12	45	29.8	17	3	R11996 N-terminal of p29 pro	1.22e+03
13	44	29.1	15	21	W15386 N-alpha-Ac-(Ala26,Phe	1.50e+03
14	44	29.1	15	20	W12816 Peptide chain ENCFE6V	1.50e+03
15	44	29.1	15	33	W51822 Peptide YX analogue #	1.50e+03

Note: Post-processor removed 968 summaries from list due to search parameters chosen.

ALIGNMENTS

RESULT ID	Score	Query Match	Length	ID	Description	Pred. No.
1	151	100.0	20	18	R92698 Human wild-type p53 gene Intron 10 encoded epitope.	5.67e-10
2	136	90.1	21	29	W33235 Human p53 fragment.	4.52e-08
3	85	56.3	20	29	W33239 Peptide of the human	7.01e-02
4	50	33.1	18	8	R44726 HEV ORF1 peptide (123	4.09e+02
5	48	31.8	17	29	W33236 Mouse p53as peptide.	6.37e+02
6	47	31.1	24	7	R37861 HIV protein fragment	7.92e+02
7	46	30.5	16	9	R45673 RNP heparin binding	9.83e+02
8	46	30.5	20	26	W26230 pMA2-TF-1g6 junction	9.83e+02
9	45	29.8	14	6	R45671 N terminus of human C	9.83e+02
10	45	29.8	14	6	R32978 Mastoparan analogue (1.22e+03
11	45	29.8	19	24	W28500 Last 19 C-terminal am	1.22e+03
12	45	29.8	17	3	R11996 N-terminal of p29 pro	1.22e+03
13	44	29.1	15	21	W15386 N-alpha-Ac-(Ala26,Phe	1.50e+03
14	44	29.1	15	20	W12816 Peptide chain ENCFE6V	1.50e+03
15	44	29.1	15	33	W51822 Peptide YX analogue #	1.50e+03

CC sequence specific binding of p53 protein is deactivated. The novel
 CC peptide differs from p53 protein in the final 50 carboxy-terminal amino
 CC acids of p53 protein, and contains a specific antigen site not present
 CC in p53 protein, giving rise to an antibody unique for the p53as peptide.
 CC Antibodies specific for murine p53as protein can be used in cell growth
 CC and differentiation basic research. The development of a homologous
 CC protein for use in human cells, would have applications in the diagnosis
 CC and prognosis of human diseases such as cancer, and in the design of
 CC treatment strategies for such diseases. The association of p53as protein
 CC expression with the G2 phase of the cell cycle suggests a functional role
 CC in G2 arrest, and the potential use of the p53as coding sequence for gene
 CC therapy.
 CC Sequence 17 AA;

Query Match 31.8%; Score 48; DB 29; Length 17;
 Best Local Similarity 55.6%; Pred. No. 6.37e+02;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 4 rafqalike 12
 :|:|:|:|
 Oy 3 RPFKALVRE 11

RESULT 6
 ID R37861 standard; peptide; 24 AA.
 AC R37861;
 DT 21-OCT-1993 (first entry)
 DE HIV protein fragment vif-C(170-192).
 KW Human immunodeficiency virus; screening; detection; AIDS;
 KW acquired immune deficiency syndrome; viral infectivity.
 OS Synthetic.
 FH Key Location/Qualifiers
 FT misc_difference 1 /note="opt.absent"
 FN J05125096-A.
 PD 21-MAY-1993.
 PR 02-NOV-1991; 288159.
 PR 02-NOV-1991; JP-288159.
 PA (YAMH) NIPPON STEEL CHEM CO.
 PA (YAMA) NIPPON STEEL CORP.
 WP: 93-200512/25.
 DT New peptide(s) - useful for diagnosis of HIV infection
 PS Claim 1; Page 2; 11pp; Japanese.
 CC This peptide has a sequence corresponding to part of the HIV
 CC C-terminal region. It is useful as a diagnostic agent for screening
 CC and monitoring HIV carriers. See also R37860.
 CC Sequence 24 AA;

Query Match 31.1%; Score 47; DB 7; Length 24;
 Best Local Similarity 60.0%; Pred. No. 7.92e+02;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Db 11 kckghrsht 20
 :|:|:|:|:|
 Oy 10 REKGHRSHS 19

RESULT 7
 ID R45673 standard; Protein; 16 AA.
 AC R45673;
 DT 25-JUL-1994 (first entry)
 DE RNP heparin binding fragment.
 KW Cationic antibacterial protein; lipopolysaccharide binding;
 KW anticoagulant; granulocytes; RNP; LPS; sepsis; autoimmune disorder;
 KW septic shock; rabbit; CAP18.
 OS Synthetic.
 PN M09402589-A.
 PD 03-FEB-1994.
 PF 15-JUL-1993; U06731.
 PR 17-JUL-1992; US-916761.
 PR 17-JUL-1992; US-916765.
 PA (PANO-) PANORAMA RES INC.
 PI Hirata M, Larrick JW, Wright SC;

DR WPI: 94-048847/06.
 PT Sequences encoding mammalian cationic antibacterial proteins -
 PT are homologous to human and rabbit CAP18 sequences and have
 PT lipopolysaccharide binding and anti-coagulation activity
 PS Disclosure; Page 50; 112pp; English.
 CC The sequence of CAP18 C-terminal RNP was compared to that of a
 CC number of heparin binding proteins to determine residues important for
 CC binding to lipopolysaccharides and inhibiting LPS-mediated activation
 CC of macrophage, as well as interfering with the clotting cascade to
 CC inhibit coagulation in conditions of disseminated intravascular
 CC coagulation.
 CC See also R45667-81.
 CC Sequence 16 AA;

Query Match 30.5%; Score 46; DB 9; Length 16;
 Best Local Similarity 45.5%; Pred. No. 9.83e+02;
 Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Db 6 lrfkrlkkek 16
 :|:|:|:|:|
 Oy 2 LRFKALVREK 12

RESULT 8
 ID W26230 standard; peptide; 20 AA.
 AC W26230;
 DT 16-MAR-1998 (first entry)
 DE pMA2-RH-19g junction region of fusion proteins.
 KW Fusion protein; hydrophilic spacer; recombinant; expression system;
 KW carboxypeptidase.
 OS Synthetic.
 PN W09728272-A1.
 PD 07-AUG-1997.
 PR 31-JAN-1997; U01470.
 PR 31-JAN-1996; US-595043.
 PA (TECH-) TECHNOLOGENE INC.
 PI Sgarlato GD;
 DR WPI: 97-402624/37.
 DR N-PSDB: R80156.
 PT Recombinant protein expression system for fusion protein production
 PT - useful for high quantity production of authentic recombinant
 PT proteins
 PS Example 1; Fig 7; 194pp; English.
 CC A novel recombinant vector has been developed which comprises a
 CC nucleotide sequence encoding a fusion protein. The fusion protein
 CC comprises three domains joined together in order, from N-terminus to
 CC C-terminus, of a first domain comprising a protein of interest, a second
 CC domain comprising a hydrophilic spacer and an affinity domain, each
 CC domain comprising amino acid residues. The present sequence represents
 CC a junction region (i.e. the region which joins the protein of interest
 CC with the affinity domain) present in pMA2-RH-19g, used in example 1
 CC of the present invention. The recombinant vector is used for the
 CC production of authentic recombinant proteins of interest. The method of
 CC the invention is useful for the expression of fusion proteins capable of
 CC isolation by affinity chromatography in pro- or eukaryotic cells. This
 CC method allows for the efficient cleavage and generation of authentic
 CC proteins of interest that do not contain extraneous (i.e. non-naturally
 CC occurring) amino acids.
 CC Sequence 20 AA;

Query Match 30.5%; Score 46; DB 26; Length 20;
 Best Local Similarity 35.3%; Pred. No. 9.83e+02;
 Matches 6; Conservative 5; Mismatches 5; Indels 1; Gaps 1;

Db 4 sfirlyp-rgrtccpc 19
 :|:|:|:|:|:|
 Oy 4 PFKALVREKGRSHSC 20

RESULT 9
 ID R45671 standard; Protein; 29 AA.
 AC R45671;
 DT 25-JUL-1994 (first entry)

DE N terminus of human CAP18.
 KM Cationic antibacterial protein; lipopolysaccharide binding;
 KM anticoagulant; granulocytes; RNP; LPS; sepsis; autoimmune disorder;
 KM septic shock; rabbit.
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT misc-difference 23 /label= Asp, Lys
 FT misc-difference 26 /label= Gln, Ile
 FT misc-difference 27 /label= Gly, Gln
 PN MO9402589-A.
 PD 03-FEB-1994.
 PE 15-JUL-1993: US-06731.
 PR 17-JUL-1992: US-916761.
 PR 17-JUL-1992: US-916765.
 PA (PANO-) PANORAMA RES INC.
 PI Hirata M, Larick JW, Wright SC;
 DR WPI: 94-048847/06.
 PT Sequences encoding mammalian cationic antibacterial proteins -
 PT are homologous to human and rabbit CAP18 sequences and have
 PT lipopolysaccharide binding and anti-coagulation activity
 PS Disclosure: Page 59; 11pp; English.
 CC The sequence is that of a human cationic antibacterial protein CAP18
 CC N-terminal fragment obtd. from granulocytes as sequenced. The
 CC sequence corresponds to the N-terminus of the C-terminal RNP fragment
 CC of rabbit CAP18. The fragment is capable of binding to lipopolysaccharide
 CC and inhibiting LPS-mediated activation of macrophage, as well as
 CC interfering with the clotting cascade to inhibit coagulation in
 CC conditions of disseminated intravascular coagulation. The polypeptides
 CC can also be used to attenuate, inhibit or prevent LPS-associated
 CC conditions, e.g. sepsis, autoimmune disorders, inflammation, etc.
 CC See also R45667-81.
 SQ Sequence 29 AA;
 Query Match 30.5%; Score 46; DB 9; Length 29;
 Best Local Similarity 45.5%; Pred. No. 9.83e+02;
 Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
 Db 6 lrfrnk16k 16
 QY 2 LRPFKALVREK 12
 RESULT 10
 ID R32978 standard; peptide: 14 AA.
 AC R32978;
 DT 02-JUL-1993 (first entry)
 DE Mastoparan analogue (41) having G protein modulatory activity.
 KM Mastoparan; MP; cellular regulation; receptor-based analogue;
 KM G protein; guanine nucleotide binding regulatory protein;
 KM asthma; ulcer; cardiovascular disease; allergy; Parkinson's;
 KM small cell carcinoma; lung; glaucoma; respiratory tract congestion;
 KM inflammation.
 OS Synthetic.
 PN MO9303748-A.
 PD 04-MAR-1993.
 PE 14-AUG-1992: US-06825.
 PR 21-AUG-1991: US-748319.
 PA (TEXA) UNIV TEXAS SYSTEM.
 PI Higashijima T, Ross EM;
 DR WPI: 93-093715/11.
 PT New peptide(s) contg. mastoparan- or receptor-analogue region -
 PT uses as G protein modulators, for treating asthma, ulcers,
 PT cardiovascular disorders and Parkinson's disease
 PS Claim 43; Page 83; 96pp; English.
 CC The peptide is an example of a highly generic formula, and is
 CC represented as found in the disclosure of the specification. The
 CC claimed peptide lacks the N-terminal 19 amino acids.
 CC The peptide is capable of modulating G protein action in a cell. It
 CC may therefore be used for treating diseases involving G proteins,
 CC e.g. asthma, ulcers, cardiovascular diseases, allergies, Parkinson's

CC disease, small cell carcinoma of the lung, glaucoma, respiratory
 CC tract congestion or inflammation.
 SQ Sequence 14 AA;
 Query Match 29.8%; Score 45; DB 6; Length 14;
 Best Local Similarity 50.0%; Pred. No. 1.22e+03;
 Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
 Db 2 nlralfar 11
 QY 1 LRPFKALVR 10
 RESULT 11
 ID W28500 standard; Protein: 19 AA.
 AC W28500;
 DT 18-NOV-1997 (first entry)
 DE Last 19 C-terminal amino acids of alternatively spliced murine p53.
 KM Leucine zipper domain; LZD; oligomerisation domain; mutant; muten;
 KM substitution; replacement; anti-oncogene; hyperproliferation;
 KM cancer; restenosis; tumour suppression; apoptosis; AS p53; mouse.
 OS Mus musculus.
 PN MO9704092-A1.
 PD 06-FEB-1997.
 PE 17-JUL-1996: F01111.
 PR 19-JUL-1995: FR-008729.
 PA (RHON) RHONE-POULENC ROBER SA.
 PI Bracco L, Conseiller E;
 DR WPI: 97-132633/12.
 DR N-P5DB; T88212.
 PT New p53 variants e.g. with oligomerisation domain replaced by
 PT leucine zipper - useful for treating hyper-proliferative disorders,
 PT esp. cancer and restenosis
 PS Example A: Page 29; 13pp; French.
 CC A claimed p53 variant consists of a fragment coding for amino
 CC acids 1-366 of human p53 protein, followed by a fragment coding for
 CC the last 19 C-terminal amino acids of the alternatively spliced (AS)
 CC form of murine p53 (i.e. the present sequence). The variant is a
 CC more active and more stable tumour suppressor and apoptosis-inducing
 CC agent than wild-type p53 and is active where the wild-type protein
 CC is not.
 SQ Sequence 19 AA;
 Query Match 29.8%; Score 45; DB 24; Length 19;
 Best Local Similarity 55.6%; Pred. No. 1.22e+03;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 Db 6 rafqajmke 14
 QY 3 RPFKALVRE 11
 RESULT 12
 ID R11996 standard; peptide: 27 AA.
 AC R11996;
 DT 26-JUL-1991 (first entry)
 DE N-terminal of p29 protein.
 KM Wegener's granulomatosis; monoclonal antibodies; autoantibodies;
 KM glomerulonephritis; serine protease; antigen.
 OS Homo sapiens.
 PN MO9106572-A.
 PD 16-MAY-1991.
 PE 29-OCT-1990: US-06827.
 PR 27-OCT-1989: US-428286.
 PA (GEHO-) GEN HOSPITAL CORP.
 PI Arnout M, McCluskey RT, Niles J;
 DR WPI: 91-164137/22.
 PT Purified p29 protein - used to detect auto-antibodies diagnostic
 PT for Wegener's granulomatosis and conditions associated with
 PT glomerulo nephritis
 PS Claim 1; Page 22; 33pp; English.
 CC The p29 protein is a 29 kD antigen which was prepd. by affinity
 CC purification. from neutrophil acid extract, using 188 monoclonal

CC antibodies. The purified antigen migrated on SDS-PAGE as three
 CC close bands, with major component at 29 kD under non-reducing
 CC conditions. It reacted with autoantibodies from patients sera
 CC indicating identity between Wegener's granulomatosis autoantigen.
 CC On isofocusing gels it had a pI of 9.2-9.4. It is a novel serine
 CC protease showing homology with leukocyte elastase and cathepsin G.
 CC The protein or monoclonal Abs can be used to test for the presence
 CC of autoantibodies diagnostic for Wegener's granulomatosis. In
 CC combination with myeloperoxidase (and/or Abs against it), p29 MAb
 CC can also be used to test for autoantibodies associated with pauci
 CC immune necrotizing and/or crescentic glomerulonephritis.
 SO Sequence 27 AA;

Query Match 29.8%; Score 45; DB 3; Length 27;
 Best Local Similarity 37.5%; Pred. No. 1.22e+03;
 Matches 6; Conservative 6; Mismatches 4; Indels 0; Gaps 0;

Db 12 RPYMASIQMGPNQSH 27
 |||||::|||
 QY 3 RPKALVREKGRPSH 18

RESULT 13
 ID W15386 standard; peptide: 15 AA.
 AC W15386:
 DT 04-JUN-1997 (first entry)
 DE N-alpha-AC-(Ala26, Phe27)-PYI(22-36).
 KM Analogue; peptide Y; PY; regulation; neuropeptide Y;
 KM intestinal water; electrolyte secretion; cell proliferation;
 KM nutrient transport; lipolysis; blood flow.
 OS Synthetic.

FH Key Location/Qualifiers
 FT modified_site 1 /note="N-alpha-acetyl-Ala"
 FT modified_site 15 /note="amidated C-terminal"
 FT modified_site 15 /note="amidated C-terminal"

PN US5604203-A.
 PD 18-FEB-1997.
 PE 29-MAR-1993; 038534.
 PR 29-MAR-1993; US-038534.
 PR 19-AUG-1993; US-109326.
 PR 24-OCT-1994; US-329151.
 PA (UICI-) UNIV CINCINNATI.
 PI Balasubramanian A;
 DR WPI; 97-144503/13.

PT New peptide YY analogues - useful for decreasing excess intestinal
 PT water and electrolyte secretion, etc.
 PS Disclosure: Column 10: 24pp; English.
 CC The sequences given in W15357-90 represent analogues of peptide YY
 CC residues 27-36 or residues 25-36. PYI shares a number of central
 CC and peripheral regulatory roles with neuropeptide Y, to which it is
 CC homologous. These peptides are used for treatment of mammals to
 CC decrease excess intestinal water and electrolyte secretion, to regulate
 CC cell proliferation, to augment nutrient transport, to regulate lipolysis
 CC and to regulate blood flow.
 SO Sequence 15 AA.

Query Match 29.1%; Score 44; DB 21; Length 15;
 Best Local Similarity 53.8%; Pred. No. 1.50e+03;
 Matches 7; Conservative 3; Mismatches 2; Indels 1; Gaps 1;

Db 2 SLRFLNLTQTR 14
 |||||::|||
 QY 1 SLRFLNLTQTR 12

RESULT 14
 ID W12816 standard; peptide: 15 AA.
 AC W12816:
 DT 21-APR-1997 (first entry)
 DE Peptide chain ENCF65VR5 of branched apogenic peptide.
 KM Branched apogenic peptide; apoptosis; inducer; erythrocyte; cancer cell;
 KM lymphocyte progenitor cell; mitogen-stimulated proliferating cell;

KM nonquiescent cell.

OS Synthetic.
 PN US5591717-A.
 PD 07-JAN-1997.
 PR 06-APR-1994; 224632.
 PR 06-APR-1994; US-224632.
 PA (CHEN/) CHENEY C M.
 PA (HART/) HARTKE J R.
 PA (ROUK/) ROUKO J L.
 PI Cheney CM, Hartke JR, Rojko JL;
 DR WPI; 97-086658/08.
 PT New branched peptide(s) which induce apoptosis - and new peptides
 PS Claim 6; Column 24; 32pp; English.
 CC W12814-W12817 represent peptides used as chains for the branched apogenic
 CC peptide of the invention. The branched apogenic peptides comprise a core
 CC sequence of at least 3, preferably at least 7 amino acids, which has at
 CC least 4 (preferably at least 8) peptide chains (such as this sequence)
 CC attached to it. These sequences were synthesised as free unattached
 CC chains, and were later joined to the core sequence. The core sequence is
 CC preferably a branched chain of lysine residues, synthesised as fully
 CC protected residues on a solid support. The peptide chain sequences are
 CC then attached to the core, before the apogenic branched peptide is
 CC deprotected and released from the solid support by reacting it with
 CC anhydrous hydrogen fluoride. The branched peptides can be used to induce
 CC apoptosis in cells in vitro, and in erythrocyte and lymphocyte progenitor
 CC cells in vivo. The peptides can also be used to induce apoptosis in
 CC cancer cells, mitogen-stimulated proliferating cells and other
 CC nonquiescent cells, to study the process and effects of apoptosis.
 SO Sequence 15 AA;

Query Match 29.1%; Score 44; DB 20; Length 15;
 Best Local Similarity 62.5%; Pred. No. 1.50e+03;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 6 QKQHGRT 13
 |||||::|||
 QY 11 EKGRPSH 18

RESULT 15
 ID W51822 standard; peptide: 15 AA.
 AC W51822:
 DT 13-OCT-1998 (first entry)
 DE Peptide YY analogue #33.
 KM Peptide YY; cell proliferation; nutrient transport; lipolysis;
 KM electrolyte secretion; anti-secretory; intestinal water; antimotility.
 OS Synthetic.
 OS Mammalia.

FH Key Location/Qualifiers
 FT Modified_site 1 /note="N-terminal acetyl"
 FT Modified_site 15 /note="C-terminal amide"
 FT Modified_site 15 /note="C-terminal amide"

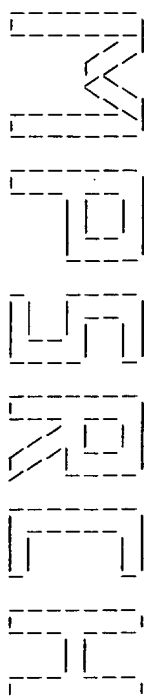
PN W09820885-A1.
 PD 22-MAY-1998.
 PE 13-NOV-1996; U18374.
 PR 13-NOV-1996; WO-U18374.
 PA (UICI-) UNIV CINCINNATI.
 PI Balasubramanian A;
 DR WPI; 98-322327/28.

PT New analogue(s) of peptide YY - used, e.g. to control cell
 PT proliferation, nutrient transport, lipolysis and intestinal water
 PT and electrolyte secretion
 PS Disclosure: Page 18; 54pp; English.
 CC The invention relates to peptide YY analogues which may be used e.g. for
 CC decreasing excess intestinal water and electrolyte secretion in mammals,
 CC to regulate cell proliferation (especially intestinal cell
 CC proliferation), to increase nutrient transport, to regulate lipolysis
 CC and to regulate blood flow. The peptides exhibit antisecretory and
 CC antimotility properties and are especially useful in treatment of
 CC gastrointestinal disorders associated with excess intestinal electrolyte
 CC and water secretion as well as decreased absorption. The new peptides

CC are truncated versions of peptide YX. They interact solely with peptide
CC YX receptors and not with homologous receptors such as NPY Y1 and Y3,
CC thus minimizing unwanted (ant)agonist side reactions. The present
CC sequence represents a peptide YX analogue.
SQ Sequence 15 AA;

Query Match 29.1%; Score 44; DB 33; Length 15;
Best Local Similarity 53.8%; Pred. No. 1.50e+03;
Matches 7; Conservative 3; Mismatches 2; Indels 1; Gaps 1;
Db 2 slraflnlvtqr 14
|||:| |||:
QY 1 SLRPFKALV-REK 12

Search completed: Wed Jun 23 14:56:08 1999
Job time : 133 secs.



(TM)

Release 3.1A John F. Collins, Biocomputing Research Unit.
Copyright (C) 1993-1998 University of Edinburgh, U.K.
Distribution rights by Oxford Molecular Ltd

MPsrch.p protein - protein database search, using Smith-Waterman algorithm

Run on: Tue Jun 22 15:00:44 1999; Maspar time 1.59 Seconds

Tabular output not generated. 128.003 Million cell updates/sec

Title: >US-08-644-289-1
Description: (1-20) from US08644289.pap
Perfect Score: 151
Sequence: 1 SLRPFKALVREKGRHSNC 20

Scoring table: PAM 150
Gap 15

Searched: 106580 seqs, 10152877 residues

Post-processing: Minimum Match 0%
Listing first 1000 summaries
Maximum DB seq length 30

Database: a-issued
1:5A_COMB 2:5B_COMB 3:PCF9_COMB 4:backfiles1

Statistics: Mean 19.456; Variance 60.473; scale 0.322

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	Pred. No.
1	136	90.1	21	1	US-08-644- Sequence 1, Applicatio	2.71e-08
2	136	90.1	21	1	US-08-259- Sequence 1, Applicatio	3.71e-08
3	85	56.3	20	1	US-08-644- Sequence 4, Applicatio	3.32e-02
4	50	33.1	18	2	US-07-876- Sequence 30, Applicatio	1.85e+02
5	48	31.8	17	1	US-08-644- Sequence 1, Applicatio	2.89e+02
6	46	30.5	16	1	US-08-313- Sequence 11, Applicati	4.48e+02
7	46	30.5	29	1	US-08-313- Sequence 7, Applicatio	4.48e+02
8	45	29.8	12	2	US-08-188- Sequence 32, Applicati	5.56e+02
9	45	29.8	14	1	US-07-603- Sequence 41, Applicatio	5.56e+02
10	45	29.8	27	1	US-07-603- Sequence 1, Applicatio	5.56e+02
11	44	29.1	15	1	US-08-224- Sequence 5, Applicatio	6.88e+02
12	44	29.1	15	1	US-08-224- Sequence 4, Applicatio	6.88e+02
13	44	29.1	20	1	US-07-882- Sequence 7, Applicatio	6.88e+02
14	43	28.5	15	1	US-08-329- Sequence 28, Applicati	8.50e+02
15	43	28.5	21	1	US-08-786- Sequence 66, Applicati	8.50e+02
16	43	28.5	21	1	US-08-786- Sequence 63, Applicati	8.50e+02
17	43	28.5	21	1	US-08-786- Sequence 67, Applicati	8.50e+02
18	43	28.5	21	1	US-08-786- Sequence 64, Applicati	8.50e+02
19	43	28.5	21	1	US-08-786- Sequence 65, Applicati	8.50e+02
20	43	28.5	22	3	US-08-485- Sequence 3, Applicatio	8.50e+02
21	43	28.5	22	1	US-08-485- Sequence 3, Applicatio	8.50e+02
22	43	28.5	23	3	PCT-US95-0 Sequence 57, Applicati	8.50e+02

23	43	28.5	23	3	PCT-US95-0	Sequence 57, Applicati	8.50e+02
24	43	28.5	24	1	US-08-786-	Sequence 60, Applicati	8.50e+02
25	43	28.5	24	1	US-08-786-	Sequence 58, Applicati	8.50e+02
26	43	28.5	24	1	US-08-786-	Sequence 56, Applicati	8.50e+02
27	43	28.5	24	1	US-08-786-	Sequence 59, Applicati	8.50e+02
28	43	28.5	27	3	PCT-US95-0	Sequence 59, Applicati	8.50e+02
29	43	28.5	27	3	PCT-US95-0	Sequence 59, Applicati	8.50e+02
30	43	28.5	27	3	PCT-US95-0	Sequence 58, Applicati	8.50e+02
31	43	28.5	27	3	PCT-US95-0	Sequence 59, Applicati	8.50e+02
32	43	28.5	28	1	US-08-786-	Sequence 42, Applicati	8.50e+02
33	43	28.5	28	1	US-08-786-	Sequence 43, Applicati	8.50e+02
34	43	28.5	28	1	US-08-786-	Sequence 45, Applicati	8.50e+02
35	43	28.5	28	1	US-08-786-	Sequence 44, Applicati	8.50e+02
36	43	28.5	28	1	US-08-786-	Sequence 46, Applicati	8.50e+02
37	42	27.8	15	1	US-08-329-	Sequence 16, Applicati	1.05e+03
38	42	27.8	15	1	US-08-329-	Sequence 17, Applicati	1.05e+03
39	42	27.8	15	1	US-08-329-	Sequence 30, Applicati	1.05e+03
40	42	27.8	15	1	US-08-329-	Sequence 17, Applicati	1.05e+03
41	42	27.8	15	1	US-08-329-	Sequence 13, Applicati	1.05e+03
42	42	27.8	15	1	US-08-329-	Sequence 27, Applicati	1.05e+03
43	42	27.8	15	1	US-08-329-	Sequence 4, Applicatio	1.05e+03
44	42	27.8	15	1	US-08-329-	Sequence 3, Applicatio	1.05e+03
45	42	27.8	16	1	US-08-488-	Sequence 33, Applicati	1.05e+03
46	42	27.8	18	3	PCT-US93-0	Sequence 2, Applicatio	1.05e+03
47	42	27.8	18	3	PCT-US93-0	Sequence 2, Applicatio	1.05e+03
48	42	27.8	24	3	PCT-US93-0	Sequence 145, Applicat	1.05e+03
49	42	27.8	24	3	PCT-US93-0	Sequence 145, Applicat	1.05e+03
50	42	27.8	24	2	US-08-488-	Sequence 145, Applicat	1.05e+03
51	42	27.8	26	2	US-08-488-	Sequence 144, Applicat	1.05e+03
52	42	27.8	26	2	US-08-488-	Sequence 144, Applicat	1.05e+03
53	42	27.8	26	3	PCT-US93-0	Sequence 144, Applicat	1.05e+03
54	41	27.2	17	3	PCT-US95-1	Sequence 42, Applicati	1.29e+03
55	41	27.2	17	2	US-08-323-	Sequence 42, Applicati	1.29e+03
56	41	27.2	17	2	US-08-413-	Sequence 62, Applicati	1.29e+03
57	41	27.2	21	2	US-08-707-	Sequence 69, Applicati	1.29e+03
58	41	27.2	21	2	US-08-477-	Sequence 97, Applicati	1.29e+03
59	41	27.2	21	2	US-08-642-	Sequence 82, Applicati	1.29e+03

Note: Post-processor removed 941 summaries from list due to search parameters chosen.

ALIGNMENTS

RESULT	ID	STANDARD:	PRT:
1	US-08-644-291-1		21 AA.
AC	xxxxxx		
DT			
XX			
DE	Sequence 1, Application US/08644291		
XX	Sequence 1, Application US/08644291		
CC	Patent No. 5726024		
CC	GENERAL INFORMATION:		
CC	APPLICANT: Kulesz-Martin, Molly F.		
CC	TITLE OF INVENTION: P53AS PROTEIN AND ANTIBODY		
CC	NUMBER OF SEQUENCES: 9		
CC	TITLE OF INVENTION: THEREFOR		
CC	CORRESPONDENCE ADDRESS:		
CC	ADDRESS: Dunn & Associates		
CC	STREET: P.O. Box 96		
CC	CITY: Newfane		
CC	STATE: New York		
CC	COUNTRY: U.S.A.		
CC	ZIP: 14108		
CC	COMPUTER READABLE FORM:		
CC	MEDIUM TYPE: Diskette - 3.50 inch, 1.44 MB		
CC	MEDIUM TYPE: Storage		
CC	COMPUTER: Victor 300 SX/25 (IBM PC Compatible)		
CC	OPERATING SYSTEM: MS-DOS Version 5.0		
CC	SOFTWARE: Wordstar Professional Release 4		
CC	CURRENT APPLICATION DATA:		


```

CC LENGTH: 20
CC TYPE: Amino Acids
CC STRANDEDNESS: unknown
CC TOPOLOGY: unknown
CC MOLECULE TYPE: Peptide
CC HYPOTHETICAL: deduced from Intron 10 sequences p53 gene
CC ANTI-SENSE:
CC FRAGMENT TYPE:
CC ORIGINAL SOURCE:
CC ORGANISM: Human
CC STRAIN:
CC INDIVIDUAL ISOLATE:
CC DEVELOPMENTAL STAGE:
CC HAPLOTYPE:
CC TISSUE TYPE:
CC CELL TYPE:
CC CELL LINE:
CC ORGANELLE:
CC IMMEDIATE SOURCE:
CC LIBRARY: deduced translation from nucleotides in
CC LIBRARY: Genbank nucleic acid database accession #54156,
CC LIBRARY: Locus HSP53G
CC CLONE:
CC POSITION IN GENOME:
CC CHROMOSOME/SEGMENT: 17
CC MAP POSITION: p53 gene, at 18530 to 18589
CC UNITS:
CC FEATURE: n/a
CC NAME/KEY:
CC LOCATION:
CC IDENTIFICATION METHOD:
CC OTHER INFORMATION:
CC PUBLICATION INFORMATION:
CC AUTHORS:
CC TITLE:
CC JOURNAL:
CC VOLUME:
CC ISSUE:
CC PAGES:
CC DATE:
CC DOCUMENT NUMBER:
CC FILING DATE:
CC PUBLICATION DATE:
CC RELEVANT RESIDUES IN SEQ ID NO:
CC SEQUENCE 20 AA; 2256 MW; 2309 CN;

SQ
Query Match 56.3%; Score 85; DB 1; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.32e-02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 REKGRRPSHC 11
| | | | | | | | | |
QY 10 REKGRRPSHC 20

RESULT 4
ID US-07-876-941A-30 STANDARD; PRT; 18 AA.
XX
XX xxxxxx
XX
XX
XX Sequence 30, Application US/07876941A
CC Sequence 30, Application US/07876941A
CC Patent No. 5885768
CC GENERAL INFORMATION:
CC APPLICANT: Reyes, Gregory R.
CC APPLICANT: Bradley, Daniel W.
CC APPLICANT: Tam, Albert W.
CC APPLICANT: Mitchell, Carl
CC TITLE OF INVENTION: Hepatitis E Virus Peptide Antigen and
CC TITLE OF INVENTION: Antibodies

```


CC PAGES: 1698-1708
CC DATE: March, 1994
CC AUTHORS: Han, K.A. and Kulesz-Martin, M.F.
CC TITLE: Alternatively Spliced p53 RNA in Transformed
CC TITLE: and No. 5747650mal Cells of Different Tissue Types
CC JOURNAL: Nucleic Acids Res.
CC VOLUME: 20
CC ISSUE: 8
CC PAGES: 1979-1981
CC DATE: 1992
CC AUTHORS: Arai, N. et al.
CC TITLE: Immunologically Distinct p53 Molecules Generated
CC TITLE: by Alternative Splicing
CC JOURNAL: Mol. and Cell. Biol.
CC VOLUME: 6
CC ISSUE: 6
CC PAGES: 3232-3239
CC DATE: 1986
CC DOCUMENT NUMBER:
CC FILING DATE:
CC PUBLICATION DATE:
CC RELEVANT RESIDUES IN SEQ ID NO:
SQ SEQUENCE 17 AA; 1944 MW; 1303 CN;

Query Match 31.8%; Score 48; DB 1; Length 17;
Best Local Similarity 55.6%; Pred. No. 2.89e+02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 4 RAFOALIKE 12
| | | | |
| | | | |
QY 3 RPFKALVRE 11

RESULT 6
ID US-08-313-681A-11 STANDARD; PRT; 16 AA.
XX xxxxxx

Sequence 11, Application US/08313681A
Patent No. 5618675
GENERAL INFORMATION:
APPLICANT: Larrick, James W.
SUSAN C.
APPLICANT: Wright, Susan C.
APPLICANT: Hirata, Mishima
TITLE OF INVENTION: Human Cationic Proteins Having
TITLE OF INVENTION: Lipopolysaccharide Binding and Anti-Coagulant Activity
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend Kourile and Crew
STREET: One Market Plaza, Stewart Tower, Suite 2000
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/313,681A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Heslin, James M.
REGISTRATION NUMBER: 29,541
REFERENCE/DOCKET NUMBER: 15325-9-1
TELEPHONE: 415-326-2400
TELECOMMUNICATION INFORMATION:

CC TELEFAX: 415-326-2422
CC INFORMATION FOR SEQ ID NO: 11:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 16 amino acids
CC TYPE: amino acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: peptide
SQ SEQUENCE 16 AA; 2071 MW; 1188 CN;

Query Match 30.5%; Score 46; DB 1; Length 16;
Best Local Similarity 45.5%; Pred. No. 4.48e+02;
Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

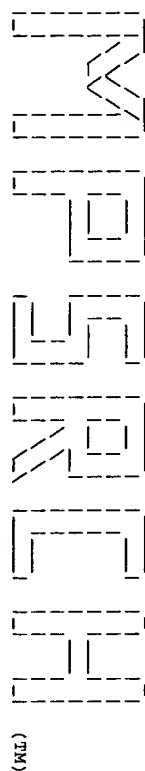
Db 6 LRFRNKIKER 16
| | | | |
| | | | |
QY 2 LRPFKALVREK 12

RESULT 7
ID US-08-313-681A-7 STANDARD; PRT; 29 AA.
XX xxxxxx

Sequence 7, Application US/08313681A
Patent No. 5618675
GENERAL INFORMATION:
APPLICANT: Larrick, James W.
SUSAN C.
APPLICANT: Wright, Susan C.
APPLICANT: Hirata, Mishima
TITLE OF INVENTION: Human Cationic Proteins Having
TITLE OF INVENTION: Lipopolysaccharide Binding and Anti-Coagulant Activity
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend Kourile and Crew
STREET: One Market Plaza, Stewart Tower, Suite 2000
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/313,681A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Heslin, James M.
REGISTRATION NUMBER: 29,541
REFERENCE/DOCKET NUMBER: 15325-9-1
TELEPHONE: 415-326-2400
TELECOMMUNICATION INFORMATION:
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 29 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Region
LOCATION: 23
OTHER INFORMATION: /note= "Xaa is Asp or Lys"
FEATURE:
NAME/KEY: Region

CC LOCATION: 26
 CC OTHER INFORMATION: /note= "Xaa is a Gln or Ile"
 CC FEATURE:
 CC NAME/KEY: Region
 CC LOCATION: 27
 CC OTHER INFORMATION: /note= "Xaa is a Gly or Gln"
 CC SEQUENCE 29 AA: 3536 MW; 5084 CN;
 SQ
 Query Match 30.5%; Score 46; DB 1; Length 29;
 Best Local Similarity 45.5%; Pred. No. 4,488+02;
 Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
 Db 6 LRRFRNKIKER 16
 |||: |||
 QY 2 LRRFKALVREK 12
 RESULT 8
 ID US-08-188-583-32 STANDARD; PRT; 12 AA.
 XX xxxxxx
 DE
 DT
 XX
 DE Sequence 32, Application US/08188583
 CC
 CC Patent No. 5851813
 CC GENERAL INFORMATION:
 CC APPLICANT: Desrosiers, Ronald C.
 CC TITLE OF INVENTION: PRIMATE LENTIVIRUS VACCINES
 CC NUMBER OF SEQUENCES: 57
 CC CORRESPONDENCE ADDRESS:
 CC ADDRESSEE: Fish & Richardson
 CC STREET: 225 Franklin Street
 CC CITY: Boston
 CC STATE: Massachusetts
 CC COUNTRY: U.S.A.
 CC ZIP: 02110-2804
 CC COMPUTER READABLE FORM:
 CC MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
 CC COMPUTER: IBM PS/2 Model 502 or 555X
 CC OPERATING SYSTEM: IBM P.C. DOS (Version 3.30)
 CC SOFTWARE: WordPerfect (Version 5.0)
 CC CURRENT APPLICATION DATA:
 CC APPLICATION NUMBER: US/08/188,583
 CC FILING DATE:
 CC CLASSIFICATION: 435
 CC PRIOR APPLICATION DATA:
 CC APPLICATION NUMBER: 07/727,494
 CC FILING DATE: July 9, 1991
 CC PRIOR APPLICATION DATA:
 CC APPLICATION NUMBER: 07/551,945
 CC FILING DATE: July 12, 1990
 CC ATTORNEY/AGENT INFORMATION:
 CC NAME: Freeman, John W.
 CC REGISTRATION NUMBER: Reg. No. 5851813 29,066
 CC TELECOMMUNICATION INFORMATION:
 CC TELEPHONE: (617) 542-5070
 CC TELEFAX: (617) 542-8906
 CC TELEX: 200154
 CC INFORMATION FOR SEQ ID NO: 32:
 CC SEQUENCE CHARACTERISTICS:
 CC LENGTH: 12
 CC TYPE: amino acid
 CC STRANDEDNESS:
 CC TOPOLOGY: linear
 CC SEQUENCE 12 AA: 1318 MW; 741 CN;
 SQ
 Query Match 29.8%; Score 45; DB 2; Length 12;
 Best Local Similarity 73.0%; Pred. No. 5,566+02;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Db 1 KGRGSHST 8
 |||||:
 QY 12 KGRPSHS 19
 RESULT 9
 ID US-08-232-453A-41 STANDARD; PRT; 14 AA.
 XX xxxxxx
 DE
 DT
 XX
 DE Sequence 41, Application US/08232453A
 CC
 CC Patent No. 589568
 CC GENERAL INFORMATION:
 CC APPLICANT: HIGASHIJIMA, TSUTOMU
 CC APPLICANT: ROSS, ELIOTT M.
 CC TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
 CC TITLE OF INVENTION: MODULATING G PROTEIN ACTION
 CC NUMBER OF SEQUENCES: 71
 CC CORRESPONDENCE ADDRESS:
 CC ADDRESSEE: ARNOLD, WHITE & DURKEE
 CC STREET: P.O. BOX 4433
 CC CITY: HOUSTON
 CC STATE: TX
 CC COUNTRY: USA
 CC ZIP: 77210
 CC COMPUTER READABLE FORM:
 CC MEDIUM TYPE: FLOPPY DISK
 CC COMPUTER: IBM PC COMPATIBLE
 CC OPERATING SYSTEM: PC-DOS/MS-DOS
 CC SOFTWARE: WORDPERFECT 5.1
 CC CURRENT APPLICATION DATA:
 CC APPLICATION NUMBER: US/08/232,453A
 CC FILING DATE: APRIL 22, 1994
 CC CLASSIFICATION: 514
 CC PRIOR APPLICATION DATA:
 CC APPLICATION NUMBER: US 07/748,319
 CC FILING DATE: AUGUST 21, 1991
 CC CLASSIFICATION: 514
 CC ATTORNEY/AGENT INFORMATION:
 CC NAME: PARKER, DAVID L.
 CC REGISTRATION NUMBER: 32,165
 CC TELECOMMUNICATION INFORMATION:
 CC TELEPHONE: (512) 418-3000
 CC TELEFAX: (512) 474-7577
 CC INFORMATION FOR SEQ ID NO: 41:
 CC SEQUENCE CHARACTERISTICS:
 CC LENGTH: 14 amino acids
 CC TYPE: amino acid
 CC STRANDEDNESS: single
 CC TOPOLOGY: linear
 CC SEQUENCE 14 AA: 1564 MW; 590 CN;
 SQ
 Query Match 29.8%; Score 45; DB 1; Length 14;
 Best Local Similarity 50.0%; Pred. No. 5,566+02;
 Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
 Db 2 NRALRALAR 11
 :|||: |||
 QY 1 SLRPFKALVR 10
 RESULT 10
 ID US-07-603-782A-1 STANDARD; PRT; 27 AA.
 XX xxxxxx
 AC
 XX
 DT



Release 3.1A John F. Collins, Biocomputing Research Unit.
Copyright (c) 1993-1998 University of Edinburgh U.K.
Distribution rights by Oxford Molecular Ltd

MPsrch_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Tue Jun 22 14:56:29 1999; Maspar time 4.30 seconds
186.302 Million cell updates/sec

Tabular output not generated.

Title: >US-08-644-289-1
Description: (1-20) from US08644289.pep
Perfect Score: 151
Sequence: 1 SLRPFKALVREKGRHPSHSC 20

Scoring table: PAM 150
Gap 15

Searched: 122810 seqs, 40068593 residues

Post-processing: Minimum Match 08
Listing first 1000 summaries
Maximum DB seq length 30

Database: p1r60
1:p1r1 2:p1r2 3:p1r3 4:p1r4

Statistics: Mean 28.488; Variance 40.935; scale 0.696

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query					
No.	Score	Match	Length	ID	Description	Pred. No.
-----	-----	-----	-----	-----	-----	-----

No matches found.

Search completed: Tue Jun 22 14:57:18 1999
Job time : 49 secs.

M P E R E H
(TM)

Release 3.1A John F. Collins, Biocomputing Research Unit.
Copyright (c) 1993-1998 University of Edinburgh, U.K.
Distribution Rights by Oxford Molecular Ltd

Msearch_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Tue Jun 22 14:57:36 1999; MasPar time 2.77 Seconds
203.787 Million cell updates/sec

Tabular output not generated.

Title: >US-08-644-289-1
Description: (1-20) from US08644289.pep
Perfect Score: 151
Sequence: 1 SLRPFKALVREKGRPSHSC 20

Scoring table:
PAM 150
Gap 15

Searched: 77977 seqs, 28268293 residues

Post-processing: Minimum Match 0%
Listing first 1000 summaries
Maximum DB seq length 30

Database: swiss-prot37
1:swissprot

Statistics: Mean 29.227; Variance 35.378; scale 0.826

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	No.	Score	Match	Length	ID	Description	Pred.	No.

No matches found.

Search completed: Tue Jun 22 14:58:30 1999
Job time : 54 secs.



Release 3.1a John F. Collins, Biocomputing Research Unit.
Copyright (c) 1993-1998 University of Edinburgh, U.K.
Distribution rights by Oxford Molecular Ltd

MProch_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Tue Jun 22 14:58:48 1999; Maspar time 5.55 Seconds

196.656 Million cell updates/sec

Tabular output not generated.

Title: >US-08-644-289-1
Description: (1-20) from US08644289.pep
Perfect Score: 151
Sequence: 1 SLRPFKALVREKGRPSHSC 20

Scoring table:
PAM 150
Gap 15

Searched: 179066 seqs, 54579741 residues

Post-processing: Minimum Match 0%
Listing first 1000 summaries
Maximum DB seq length 30

Database: splrembl9
1:sp_archaea 2:sp_bacteria 3:sp_fungi 4:sp_human
5:sp_invertebrate 6:sp_mammal 7:sp_mhc 8:sp_organelle
9:sp_phase 10:sp_plant 11:sp_rodent 12:sp_unclassified
13:sp_vertebrate 14:sp_virus

Statistics: Mean 28.060; Variance 36.106; scale 0.777

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match	Length	ID	Description	Pred. No.
------------	-------------	--------	----	-------------	-----------

No matches found.

Search completed: Tue Jun 22 15:00:25 1999
Job time : 97 secs.